In the Claims

Please cancel claims 14-15 without prejudice to applicants' right to pursue the subject matter of these claims in this or a subsequent application.

1. - 11. (Canceled)

12.(Previously Presented): N-(3-ethynylphenyl)-6,7-bis(2-methoxyethoxy)-4-quinazolinamine mesylate.

13.(Previously Presented): A hydrate form of N-(3-ethynylphenyl)-6,7-bis(2-methoxyethoxy)-4-quinazolinamine mesylate of claim 12.

14 - 15.(Cancelled)

16.(Currently Amended): ~~The~~ A pharmaceutical composition ~~of claim 15,~~ comprising from about 0.001 mg to about 100 mg of ~~the compound~~ N-(ethynylphenyl)-6,7-bis(2-methoxyethoxy)-4-quinazolinamine mesylate and a pharmaceutically acceptable carrier.

17.(Previously Presented): The pharmaceutical composition of claim 16, comprising from about 1 mg to about 35 mg of the compound.

18.(Previously Presented): The pharmaceutical composition of claim 16, comprising from about 0.05 mg to about 7 g of the compound.

19.(Previously Presented): The pharmaceutical composition of claim 17, comprising from about 0.2 g to about 2.5 g of the compound.

20.(Previously Presented): The pharmaceutical composition of claim 19, in the form of a tablet, capsule, pill, powder, sustained release formulations, solution, parenteral injection as a sterile solution, suspension or emulsion, or suppository.

21.(Previously Presented): The pharmaceutical composition of claim 20, in the form of a parenteral injection.

22.(Previously Presented): The pharmaceutical composition of claim 20, in the form of a tablet.

23.(Currently Amended): A method of treating a mammal suffering from a hyperproliferative disorder which comprises administering to said mammal an amount of the ~~compound~~ pharmaceutical composition of claim ~~12~~ 16 therapeutically effective to inhibit the epidermal growth factor receptor ("EGFR") in the mammal, so as to thereby treat the mammal.

24.(Previously Presented): The method of claim 23 wherein the hyperproliferative disorder is a cancer selected from the group consisting of brain, lung, squamous cell, bladder, gastric, pancreatic, breast, head, neck, renal, kidney, ovarian, prostate, colorectal, oesophageal, gynecological and thyroid cancer.

25.(Currently Amended): The method of claim 23 further comprising administering to said mammal a therapeutically effective amount of a compound selected from the group consisting of ~~mitotic inhibitors,~~ alkylating agents, anti-metabolites, intercalating antibiotics, growth factor inhibitors, cell-cycle inhibitor, enzymes, topoisomerase, inhibitors, ~~biological response modifiers,~~ anti-hormones, and anti-androgens.

26. (New): The method of claim 25 wherein the cell-cycle inhibitor is a mitotic inhibitor.